Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application.

- 1. (Currently Amended) A compound comprising:
- (a) a CD1d complex; and
- (b) an antibody specific for a cell surface marker or an antigen-binding fragment thereof;

wherein said CD1d complex comprises a CD1d molecule or fragment thereof, and a β2-microglobulin molecule or fragment thereof, and a lipid or glycolipid antigen associated with said CD1d molecule; and wherein said CD1d complex is linked to said antibody or fragment thereof.

- 2. (Canceled)
- 3. (Canceled)
- 4. (Currently Amended) The compound of claim $3\ \underline{1}$, wherein said antigen is α -GalCer.
- 5. (Withdrawn, Currently Amended) The compound of claim 3 1, wherein said antigen is α -GalCer modified to have a shortened long-chain sphingosine base (C5 vs. C14) and acyl chain (C24 vs. C26)
- 6. (Withdrawn) The compound of claim 5, wherein said modified α-GalCer is the OCH analog with a long-chain sphingosine base shortened from C14 to C5 and acyl chain from C26 to C24.
- 7. (Withdrawn) The compound of claim 1, wherein said antibody fragment is a F(ab).

- 8. (Previously Presented) The compound of claim 1, wherein said antibody fragment is a scFv.
- 9. (Withdrawn) The compound of claim 1, wherein said antibody is a full-length antibody.
- 10. (Currently Amended) The compound of any of claims 1 or 4-9, wherein said cell surface marker is a cell surface marker of tumor cells.
- 11. (Original) The compound of claim 10, wherein said cell surface marker is selected from the group consisting of: CEA, Her2/neu, EGFR type I or type II, CD19, CD20, CD22, Muc-1, PSMA, or STEAP.
- 12. (Canceled)
- 13. (Withdrawn) The compound of claim 1, wherein said cell surface marker is a cell surface marker of dendritic cells.
- 14-16. (Canceled)
- 17. (Withdrawn) The compound of claim 1, wherein said cell surface marker is a cell surface marker of a target of autoimmune or inflammatory disease.
- 18-27. (Canceled)
- 28. (Withdrawn) The compound of claim 1, wherein said cell surface marker is a cell surface marker of an infected cell or tissue.
- 29-35. (Canceled)
- 36. (Withdrawn) The compound of claim 1, wherein said CD1d molecule is attached to the heavy chain of said antibody.
- 37. (Withdrawn) The compound of claim 1, wherein said CD1d molecule is attached to the light chain of said antibody.

- 38. (Withdrawn) The compound of claim 1, wherein said β 2 microglobulin molecule is attached to the heavy chain of said antibody.
- 39. (Withdrawn) The compound of claim 1, wherein said β 2 microglobulin molecule is attached to the light chain of said antibody.
- 40. (Previously Presented) The compound of claim 1, wherein the CD1d complex is linked in a fusion protein with the antibody or fragment thereof.
- 41. (Withdrawn) The compound of claim 1, wherein said CD1d complexes are attached to said antibody through a linker sequence.
- 42-45. (Canceled)
- 46. (Withdrawn) A method of inducing an anti-tumor response in a mammal, comprising administering the compound of claim 49 to said mammal.
- 47-48. (Canceled)
- 49. (Previously Presented) The compound of claim 11, wherein said cell surface marker is Her2/neu.
- 50. (Previously Presented) The compound of claim 1, wherein the CD1d molecule or fragment thereof of said CD1d complex comprises the extracellular portion of CD1d.
- 51. (Previously Presented) The compound of claim 50, wherein said extracellular portion comprises amino acids 1-297 of the amino acid sequence of SEQ ID NO: 40.
- 52. (Previously Presented) The compound of claim 8, wherein the variable light domain and the variable heavy domain of said scFv are linked by a peptide bridge.
- 53. (Withdrawn) The compound of claim 8, wherein the variable light domain and the variable heavy domain of said scFv are linked by one or more disulfide bonds.

- 54. (Previously Presented) The compound of claim 40, wherein the CD1d molecule of said CD1d complex is fused to said antibody or fragment thereof.
- 55. (Previously Presented) The compound of claim 54, wherein the CD1d molecule of said CD1d complex is linked to the amino terminus of the antibody or fragment thereof.
- 56. (Withdrawn) The compound of claim 54, wherein the CD1d molecule of said CD1d complex is linked to the carboxyl terminus of the antibody or fragment thereof.
- 57. (Previously Presented) The compound of claim 54, wherein a short linker amino acid sequence of from about 3 to about 30 amino acids is situated between the CD1d molecule of said CD1d complex and the antibody or fragment thereof.
- 58. (Withdrawn) The compound of claim 57, wherein said short linker amino acid sequence comprises the sequence of SEQ ID NO: 1.
- 59. (Previously Presented) The compound of claim 57, wherein said short linker amino acid sequence comprises the sequence of SEQ ID NO: 2.
- 60. (Withdrawn) The compound of claim 40, wherein the β 2-microglobulin molecule of said CD1d complex is fused to said antibody or fragment thereof.
- 61. (Withdrawn) The compound of claim 60, wherein the β 2-microglobulin molecule of said CD1d complex is linked to the amino terminus of the antibody or fragment thereof.
- 62. (Withdrawn) The compound of claim 60, wherein the β 2-microglobulin molecule of said CD1d complex is linked to the carboxyl terminus of the antibody or fragment thereof.
- 63. (Withdrawn) The compound of claim 60, wherein a short linker amino acid sequence of from about 3 to about 30 amino acids is situated between the β 2-microglobulin molecule of said CD1d complex and the antibody or fragment thereof.

- 64. (Withdrawn) The compound of claim 63, wherein said short linker amino acid sequence comprises the sequence of SEQ ID NO: 1.
- 65. (Withdrawn) The compound of claim 63, wherein said short linker amino acid sequence comprises the sequence of SEQ ID NO: 2.